

AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims

Claim 1 (currently amended): A method ~~for~~ of producing a transgenic cotton plant comprising the steps of:

- (a) obtaining cotton petiole explants,
- (b) exposing the petiole explants to a culture of *Agrobacterium tumefaciens* that harbors a vector comprising an exogenous gene and a selectable marker gene in medium that does not contain plant hormones and contains glucose as the sole carbon source, the *Agrobacterium* being capable of effecting the stable transfer of the exogenous gene and selectable marker gene to the genome of the cells of the petiole explant,
- (c) culturing the petiole explants ~~in~~ on medium containing one or more plant hormones and contains glucose as the sole carbon source to induce callus formation, wherein the one or more plant hormones is 2,4-dichlorophenoxyacetic acid at a concentration up to about 0.5 mg/l and kinetin at a concentration up to about 1 mg/l,
- (d) selecting a transformed callus that expresses the exogenous gene on medium that does not contain plant hormones and contains glucose as the sole carbon source,
- (e) culturing the selected callus in suspension culture in medium that does not contain plant hormones and contains glucose as the sole carbon source for a duration of less than about 20 days to induce formation of embryogenic calli,
- (f) culturing the embryogenic calli on medium that does not contain plant hormones and contains glucose as the sole carbon source to induce formation of embryoids, and
- (g) germinating an embryoid on a medium that does not contain plant hormones, contains glucose as the sole carbon source and contains a source of nitrogen selected from the group

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consisting of asparagine, glutamine and both asparagine and glutamine to obtain a young transgenic cotton plant.

Claim 2 (previously presented): The method of claim 1, wherein the petiole explants are pre-cultured for a period of time prior to exposure to the culture of *Agrobacterium tumefaciens*.

Claim 3 (canceled).

Claim 4 (currently amended): The method of claim 3 1, wherein the glucose is at a concentration of about 10 g/l to about 50 g/l.

Claim 5 (previously presented): The method of claim 4, wherein the glucose is at a concentration of about 30 g/l.

Claims 6-7 (canceled).

Claim 8 (currently amended): The method of claim 7 1, wherein the source of nitrogen is at a concentration of about 700 mg/l to about 5 g/l.

Claim 9 (currently amended): The method of claim 7 1, wherein the medium further contains KNO₃ as a further source of nitrogen at a concentration of about 3.8 g/l.

Claim 10 (currently amended): The method of claim 7 1, wherein the source of nitrogen is both asparagine and glutamine, and the asparagine is at a concentration of about 200 mg/l to about 1 g/l and the glutamine is at a concentration of about 500 mg/l to about 2 g/l.

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Claim 11 (previously presented): The method of claim 10, wherein the asparagine is at a concentration of about 500 mg/l and the glutamine is at a concentration of about 1 g/l.

Claim 12 (canceled).

Claim 13 (previously presented): The method of claim 1, wherein the suspension culture of step (e) has a duration of about 10 days to about 20 days.

Claim 14 (previously presented): The method of claim 13, wherein the suspension culture of step (e) has a duration of about 14 days.

Claims 15-17 (canceled).

Claim 18 (previously presented): The method of claim 1, wherein the 2,4-dichlorophenoxyacetic acid is at a concentration of about 0.05 mg/l and the kinetin is at a concentration of about 0.1 mg/l.

Claim 19 (currently amended): A method ~~for of~~ producing a transgenic cotton plant comprising the steps of:

- (a) obtaining tender petiole explants from cotton plants,
- (b) exposing the petiole explants to a culture of *Agrobacterium tumefaciens* that harbors a vector comprising an exogenous gene and a selectable marker gene in a medium that does not contain plant hormones and contains glucose as the sole carbon source, the *Agrobacterium* being capable of effecting the stable transfer of the exogenous gene and selectable marker gene to the genome of the cells of the petiole explant,

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- (c) culturing the petiole explants to induce callus formation ~~in~~ on medium containing about 0.05 mg/l 2, 4-dichlorophenoxyacetic acid and about 0.1 mg/l kinetin and glucose as the sole carbon source,
- (d) selecting a transformed callus that expresses the exogenous gene on medium that does not contain plant hormones and contains glucose as the sole carbon source,
- (e) culturing the selected callus in suspension culture in medium that does not contain containing no added plant hormones and contains glucose as the sole carbon source for a duration of less than about 20 days to induce formation of embryogenic calli,
- (f) culturing the embryogenic calli on medium that does not contain plant hormones and contains glucose as the sole carbon source to induce formation of embryooids, and
- (g) germinating an embryooid on medium that does not contain plant hormones, contains glucose as the sole carbon source, contains KNO₃ at a concentration of 3.8 g/l and contains a further source of nitrogen selected from the group consisting of asparagine, glutamine and both asparagine and glutamine to obtain a young transgenic cotton plant ~~on a medium containing KNO₃ at a concentration of 3.8 g/l~~.

Claim 20 (currently amended): The method of claim 1 which further comprises:

- (h) growing the young transgenic cotton plant on a medium that does not contain plant hormones and contains glucose and sucrose as the carbon source to produce a transgenic cotton plant capable of growth in soil.

Claim 21 (canceled).

Claim 22 (currently amended): The method of claim ~~2+~~ 20, wherein the medium contains about 10 g/l of each of the glucose and the sucrose.

Claim 23 (currently amended): The method of claim 19 which further comprises:

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(h) growing the young transgenic cotton plant on a medium that does not contain plant hormones and contains glucose and sucrose as the carbon source to produce a transgenic cotton plant capable of growth in soil.

Claim 24 (canceled).

Claim 25 (currently amended): The method of claim 24 23, wherein the medium contains about 10 g/l of each of the glucose and the sucrose.

Claim 26 (canceled)

Claim 27 (currently amended): The method of claim 26 23, wherein the asparagine is at a concentration of about 500 mg/l and the glutamine is at a concentration of about 1 g/l.

Claim 28 (canceled).

Claim 29 (currently amended): The method of claim 28 19, wherein the suspension culture of step (e) has a duration of about 10 days to about 20 days.

Claim 30 (currently amended): The method of claim 28 19, wherein the suspension culture of step (e) has a duration of about 14 days.

Claim 31 (new): The method of claim 1, wherein the pH of the media in steps (c)-(f) is 6.2 to 7.0.

Claim 32 (new): The method of claim 1, wherein the pH of the media in steps (c)-(g) is 6.5.

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Claim 33 (new): The method of claim 19, wherein the pH of the media in steps (c)-(f) is 6.2 to 7.0.

Claim 34 (new): The method of claim 19, wherein the pH of the media in steps (c)-(g) is 6.5.

Claim 35 (new): The method of claim 20, wherein the pH of the medium in step (h) is 7.0.

Claim 36 (new): The method of claim 23, wherein the pH of the medium in step (h) is 7.0.